nature portfolio

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Reporting Summary

Nature Portfolio wishes to improve the reproducibility of the work that we publish. This form provides structure for consistency and transparency in reporting. For further information on Nature Portfolio policies, see our <u>Editorial Policies</u> and the <u>Editorial Policy Checklist</u>.

Statistics					
	ses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.				
n/a Confirmed	,,,,,,,,,,,,,,,,,,,,				
The exact sam	mple size (n) for each experimental group/condition, given as a discrete number and unit of measurement				
	on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly				
The statistical Only common t	The statistical test(s) used AND whether they are one- or two-sided Only common tests should be described solely by name; describe more complex techniques in the Methods section.				
	of all covariates tested				
A description	of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons				
A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)					
For null hypothesis testing, the test statistic (e.g. <i>F</i> , <i>t</i> , <i>r</i>) with confidence intervals, effect sizes, degrees of freedom and <i>P</i> value noted Give <i>P</i> values as exact values whenever suitable.					
For Bayesian	analysis, information on the choice of priors and Markov chain Monte Carlo settings				
For hierarchic	cal and complex designs, identification of the appropriate level for tests and full reporting of outcomes				
Estimates of e	effect sizes (e.g. Cohen's d , Pearson's r), indicating how they were calculated				
'	Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.				
Software and o	code				
Policy information abo	out <u>availability of computer code</u>				
Data collection Qu	uantstudio™ Design & Analysis Software v1.52				
Data analysis Qu	uantstudio™ Design & Analysis Software v1.52, GraphPad Prism 9				
For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors and reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Portfolio guidelines for submitting code & software for further information.					
Data					
Policy information abo	· · · · · · · · · · · · · · · · · · ·				
- Accession codes, un	include a <u>data availability statement</u> . This statement should provide the following information, where applicable: nique identifiers, or web links for publicly available datasets y restrictions on data availability				

- For clinical datasets or third party data, please ensure that the statement adheres to our policy

All data upon which conclusions are made are included in the manuscript or in the supplemental information file provided.

Research involving human participants, their data, or biological material

Policy information about studies with human participants or human data. See also policy information about sex, gender (identity/presentation), and sexual orientation and race, ethnicity and racism.

Reporting on sex and gender

The findings are not limited to only one biologic sex. Biologic sex was not considered in the study design. Sex refers to the sex assigned at birth which in all cases was confirmed at autopsy by examination of internal and external organs. The requirement for patient consent for this project was waived by the IRB, and patient consent was not obtained. The study involved a total of 15 male decedents and 10 female decedents. The detection of vaccine in the heart was not associated with sex.

Reporting on race, ethnicity, or other socially relevant groupings

Race was categorized as white, black, or unknown based on the medical record. Confounding variables regarding race were not controlled for due to the small sample size.

Population characteristics

The vaccinated patients entailed 20 patients with a mean age of 64, a mean BMI of 26 and histories of coronary artery disease (33%), hypertension (78%), hyperlipidemia (72%), diabetes mellitus (33%), autoimmune disease (17%), and smoking

Recruitment

The patients recruited all underwent autopsy performed at Massachusetts General Hospital between January 2021 and February 2022. Inclusion criteria were the ability to collect fresh tissue, a clear history of either vaccination or no vaccination, and a post-mortem interval ≤ 60 hours. Patients with a history of SARS-CoV-2 infection were excluded.

Ethics oversight

Mass General Brigham IRB

Note that full information on the approval of the study protocol must also be provided in the manuscript.

Field-specific reporting

Please select the one belor	w that is the best fit for your research	n. If you are not sure, read the appropriate sections before making your selection.			
X Life sciences	Behavioural & social sciences	Ecological, evolutionary & environmental sciences			
or a reference copy of the document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>					
	and the design				
Life sciences	s study design				

All studies must disclose on these points even when the disclosure is negative.

Sample size	Samples size was determined based on the number of autopsies that met the study criteria.	
Data exclusions	No data were excluded.	
Replication	All PCR-positive samples were confirmed by sequencing.	
Randomization	Samples were allocated into groups based on the presence or absence of vaccine detected by RT-PCR. The number samples positive for vaccine were small and covariates were not controlled.	
Blinding	The investigator performing the vaccine RT-PCR was blinded to the presence of myocardial injury and the macrophage counts.	

Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

Materials & experime	ental systems Methods	
n/a Involved in the study	n/a Involved in the study	
Antibodies	ChIP-seq	
Eukaryotic cell lines	rs Flow cytometry	
Palaeontology and a	l archaeology MRI-based neuroimaging	
Animals and other o	organisms	
Clinical data		
Dual use research of	of concern	
Plants		
Antibodies		
Antibodies used	SARS-CoV-2 Spike Protein S1 rabbit Polyclonal Antibody, Invitrogen, Waltham, MA, catalogue # PA5-114528 Anti-CD68 Monoclonal Antibody, clone KP1, eBioscienceTM, Life Technologies, Carlsbad, CA, catalogue # 14-06	88-82
Validation	Lung tissue from patients who died of severe COVID-19 was used as a positive control for both antibodies.	
Clinical data		
Policy information about cli	clinical studies	
All manuscripts should comply	ly with the ICMJE <u>guidelines for publication of clinical research</u> and a completed <u>CONSORT checklist</u> must be includ	ed with all submissions.
Clinical trial registration	Not applicable, this was not a clinical trial.	
Study protocol	Not applicable, this was not a clinical trial	
Data collection	The patients recruited all underwent autopsy performed at Massachusetts General Hospital between January 2 2022. Clinical data was collected following the autopsies.	2021 and February

Study outcomes were the presence or absence of vaccine in the tissues as determined by RT-PCR.

Outcomes